

10/518,007

=> d his

(FILE 'HOME' ENTERED AT 13:49:37 ON 16 NOV 2006)

FILE 'REGISTRY' ENTERED AT 13:49:58 ON 16 NOV 2006

L1 STRUCTURE UPLOADED

L2 2 S L1

L3 8 S L1 FULL

FILE 'HCAPLUS, CHEMCATS' ENTERED AT 13:50:50 ON 16 NOV 2006

L4 2 S L3

FILE 'HCAPLUS, HCAOLD, USPATFULL, EPFULL, BIOSIS, MEDLINE' ENTERED AT
13:51:47 ON 16 NOV 2006

L5 188036 S BENZOIC ACID? OR BENZOIC ACID DERIVATIVE?

L6 103714 S L5 AND ?AMINO?

L7 680313 S DIABETES OR INSULIN RESISTANCE OR PEROXISOME PROLIFERATOR-ACT

L8 10162 S L6 AND L7

L9 4662 S L8 AND LIPID?

L10 4647 S L9 AND (PHARMACEUTICAL OR TREAT? OR COMPOSITION)

L11 4406 S L10 AND (SALT OR PRODRUG)

L12 3284 S L11 AND (OXY OR SULFUR OR SULFONYL)

L13 901 S L12 AND ORTHO

L14 508 S L13 AND (PHENOXY OR OXOETHYL OR OXOPROPYL)

L15 487 S L14 AND BENZYL

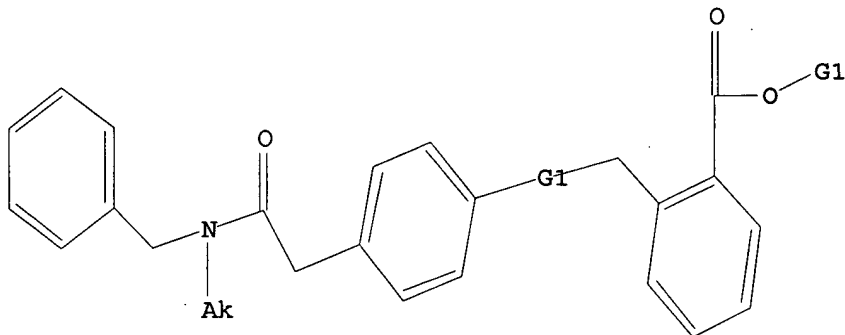
10/518,007

L1 STRUCTURE UPLOADED

=> d

L1 HAS NO ANSWERS

L1 STR



G1 X, Cb, Cy, Hy, Ak

Structure attributes must be viewed using STN Express query preparation.

=> s l1

SAMPLE SEARCH INITIATED 15:23:43 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 138 TO ITERATE

100.0% PROCESSED 138 ITERATIONS

0 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS: 2056 TO 3464

PROJECTED ANSWERS: 0 TO 0

L2 0 SEA SSS SAM L1

=> s l1 full

FULL SEARCH INITIATED 15:23:46 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 3141 TO ITERATE

100.0% PROCESSED 3141 ITERATIONS

0 ANSWERS

SEARCH TIME: 00.00.01

L3 0 SEA SSS FUL L1

=>

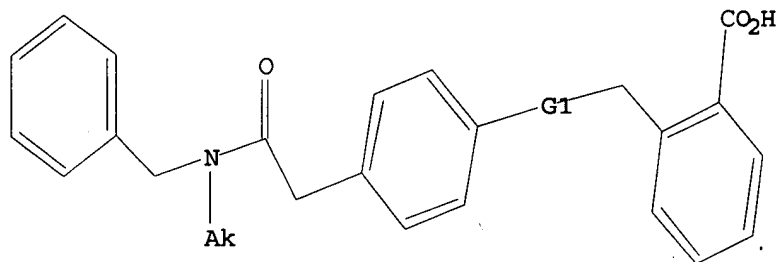
10/518,007

L1 STRUCTURE UPLOADED

=> d

L1 HAS NO ANSWERS

L1 STR



G1 O,S

Structure attributes must be viewed using STN Express query preparation.

=> s l1

SAMPLE SEARCH INITIATED 13:50:17 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 8 TO ITERATE

100.0% PROCESSED
SEARCH TIME: 00.00.01

8 ITERATIONS

2 ANSWERS

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**

PROJECTED ITERATIONS: 8 TO 329

PROJECTED ANSWERS: 2 TO 124

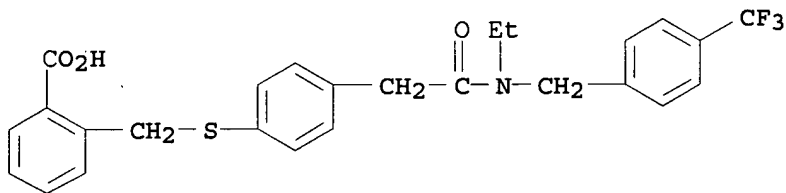
L2 2 SEA SSS SAM L1

=> d scan

L2 2 ANSWERS REGISTRY COPYRIGHT 2006 ACS on STN

IN Benzoic acid, 2-[[[4-[2-[ethyl[[4-(trifluoromethyl)phenyl]methyl]amino]-2-oxoethyl]phenyl]thio]methyl]- (9CI)

MF C26 H24 F3 N O3 S



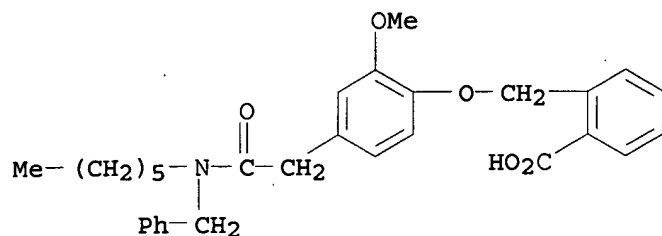
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):1

L2 2 ANSWERS REGISTRY COPYRIGHT 2006 ACS on STN

IN Benzoic acid, 2-[[[4-[2-[hexyl(phenylmethyl)amino]-2-oxoethyl]-2-

methoxyphenoxy]methyl] - (9CI)
MF C30 H35 N O5



ALL ANSWERS HAVE BEEN SCANNED

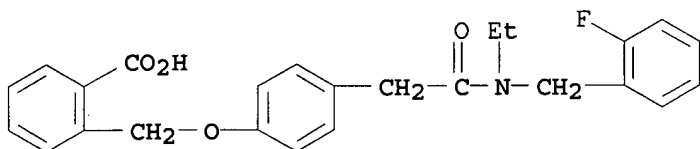
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=> s ll full
FULL SEARCH INITIATED 13:50:33 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED -      74 TO ITERATE
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100.0% PROCESSED          74 ITERATIONS                      8 ANSWERS
SEARCH TIME: 00.00.01
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L3 8 SEA SSS FUL L1

=> d scan

L3 8 ANSWERS REGISTRY COPYRIGHT 2006 ACS on STN
IN Benzoic acid, 2-[[4-[2-[ethyl[(2-fluorophenyl)methyl]amino]-2-
oxoethyl]phenoxy]methyl]- (9CI)
MF C25 H24 F N O4

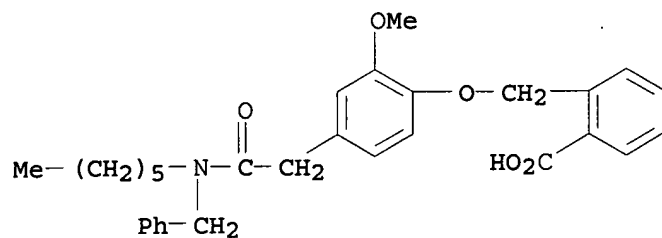


PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

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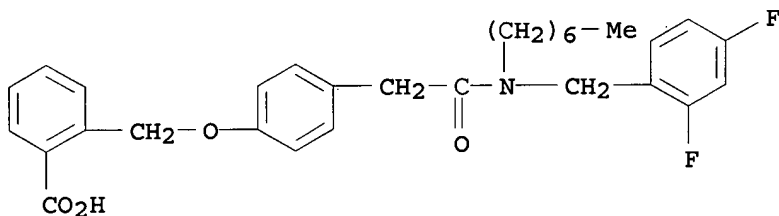
L3 8 ANSWERS REGISTRY COPYRIGHT 2006 ACS on STN
IN Benzoic acid, 2-[[4-[2-[hexyl(phenylmethyl)amino]-2-oxoethyl]-2-
methoxyphenoxy]methyl]- (9CI)
MF C30 H35 N O5

10/518,007



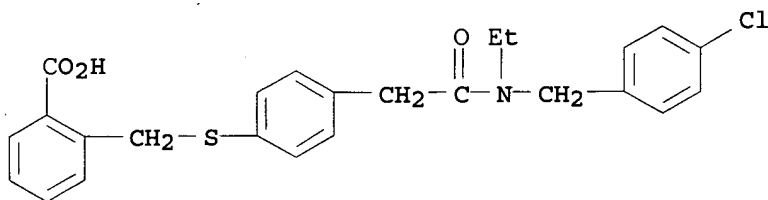
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L3 8 ANSWERS REGISTRY COPYRIGHT 2006 ACS on STN
IN Benzoic acid, 2-[[4-[2-[[[(2,4-difluorophenyl)methyl]heptylamino]-2-oxoethyl]phenoxy]methyl]- (9CI)
MF C30 H33 F2 N O4



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

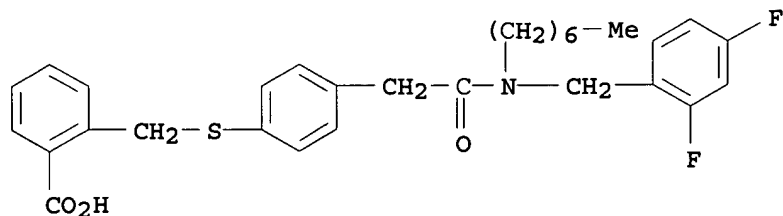
L3 8 ANSWERS REGISTRY COPYRIGHT 2006 ACS on STN
IN Benzoic acid, 2-[[[4-[2-[[[(4-chlorophenyl)methyl]ethylamino]-2-oxoethyl]phenyl]thio]methyl]- (9CI)
MF C25 H24 Cl N O3 S



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

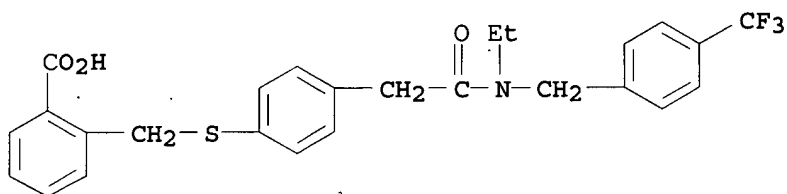
L3 8 ANSWERS REGISTRY COPYRIGHT 2006 ACS on STN
IN Benzoic acid, 2-[[[4-[2-[[[(2,4-difluorophenyl)methyl]heptylamino]-2-oxoethyl]phenyl]thio]methyl]- (9CI)
MF C30 H33 F2 N O3 S

10/518,007



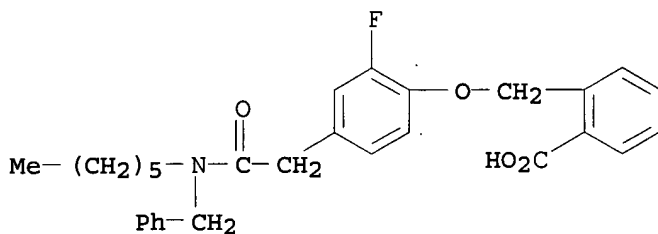
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L3 8 ANSWERS REGISTRY COPYRIGHT 2006 ACS on STN
IN Benzoic acid, 2-[[[4-[2-[ethyl[[4-(trifluoromethyl)phenyl]methyl]amino]-2-oxoethyl]phenyl]thio]methyl]- (9CI)
MF C26 H24 F3 N O3 S



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

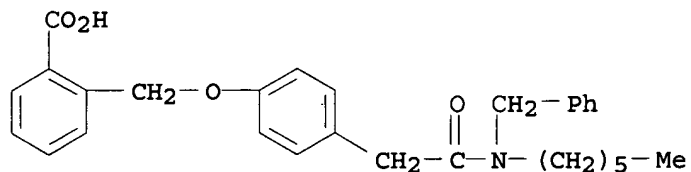
L3 8 ANSWERS REGISTRY COPYRIGHT 2006 ACS on STN
IN Benzoic acid, 2-[[[4-[2-[hexyl(phenylmethyl)amino]-2-oxoethyl]phenoxy]methyl]- (9CI)
MF C29 H32 F N O4



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L3 8 ANSWERS REGISTRY COPYRIGHT 2006 ACS on STN
IN Benzoic acid, 2-[[[4-[2-[hexyl(phenylmethyl)amino]-2-oxoethyl]phenoxy]methyl]- (9CI)
MF C29 H33 N O4

10/518,007



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

ALL ANSWERS HAVE BEEN SCANNED

=> file hcaplus chemcat

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

FULL ESTIMATED COST

166.94

167.15

FILE 'HCAPLUS' ENTERED AT 13:50:50 ON 16 NOV 2006

USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.

PLEASE SEE "HELP USAGETERMS" FOR DETAILS.

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FILE 'CHEMCATS' ENTERED AT 13:50:50 ON 16 NOV 2006

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=> s l3

L4 2 L3

=> d 1-2 ibib abs hitstr

L4 ANSWER 1 OF 2 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2004:2838 HCAPLUS

DOCUMENT NUMBER: 140:41910

TITLE: Preparation of ortho-substituted benzoic acid derivatives for the treatment of insulin resistance

INVENTOR(S): Li, Lanna

PATENT ASSIGNEE(S): Astrazeneca Ab, Swed.

SOURCE: PCT Int. Appl., 60 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

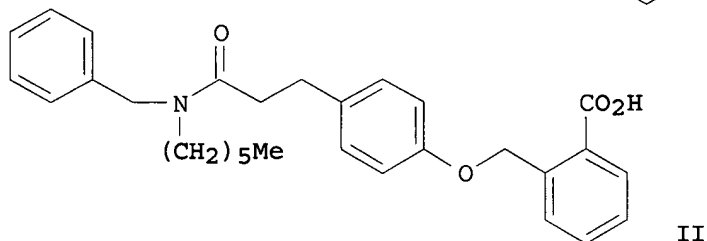
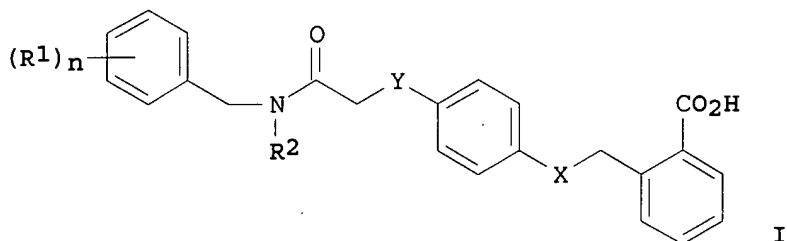
FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

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W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			

CA 2489834	AA	20031231	CA 2003-2489834	20030617
CA 2489834	C	20061003		
AU 2003240099	A1	20040106	AU 2003-240099	20030617
BR 2003011932	A	20050322	BR 2003-11932	20030617
EP 1517883	A1	20050330	EP 2003-732713	20030617
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CN 1675171	A	20050928	CN 2003-819474	20030617
JP 2005529970	T2	20051006	JP 2004-515007	20030617
JP 3782818	B2	20060607		
NO 2004005353	A	20050120	NO 2004-5353	20041207
US 2005256198	A1	20051117	US 2004-518007	20041214
ZA 2004010161	A	20051020	ZA 2004-10161	20041215
JP 2006182782	A2	20060713	JP 2006-19068	20060127
PRIORITY APPLN. INFO.:			SE 2002-1935	A 20020620
			SE 2002-3826	A 20021220
			JP 2004-515007	A3 20030617
			WO 2003-GB2584	W 20030617

OTHER SOURCE(S): MARPAT 140:41910
GI



AB Title compds. I [$n = 0-2$; $R_1 = \text{halo, alkyl, alkoxy, etc.}$; $R_2 = \text{alkyl}$; $Y = \text{absent, CH}_2$; $X = O, S$] are prepared. For instance, N-benzyl-N-hexyl-3-(4-hydroxyphenyl)propanamide (preparation given) is reacted with Me 2-(bromomethyl)benzoate (CH_3CN , K_2CO_3 , 66°) and the product saponified ($\text{THF}/\text{H}_2\text{O}$, LiOH , microwave, 120° , 40 min) to give II. Example compds. have $\text{EC}_{50} < 50 \mu\text{mol/L}$ for PPAR- α . I are useful for treating clin. conditions associated with insulin resistance.

IT 637015-02-2P, 2-[[[4-[2-[(2,4-Difluorobenzyl) (heptyl)amino]-2-oxoethyl]phenyl]thio]methyl]benzoic acid 637015-05-5P, 2-[[[4-[2-[Benzyl (hexyl)amino]-2-oxoethyl]phenoxy]methyl]benzoic acid 637015-07-7P, 2-[[[4-[2-[(2,4-Difluorobenzyl) (heptyl)amino]-2-oxoethyl]phenoxy]methyl]benzoic acid 637015-32-8P, 2-[[[4-[2-[Ethyl[4-(trifluoromethyl)benzyl]amino]-2-oxoethyl]phenyl]thio]methyl]benzoic acid 637015-33-9P, 2-[[[4-[2-[Ethyl(2-fluorobenzyl)amino]-2-oxoethyl]phenoxy]methyl]benzoic acid 637015-40-8P, 2-[[[4-[2-[(4-Chlorobenzyl) (ethyl)amino]-2-oxoethyl]phenyl]thio]methyl]benzoic acid

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU

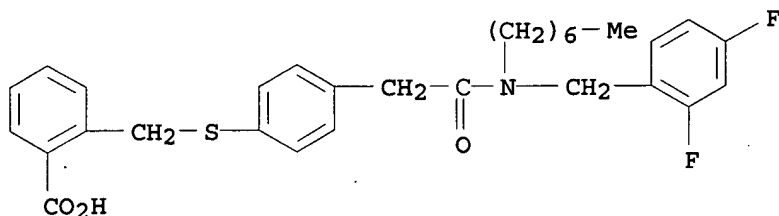
10/518,007

(Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(ortho-substituted benzoic acid derivs. for treatment of insulin resistance)

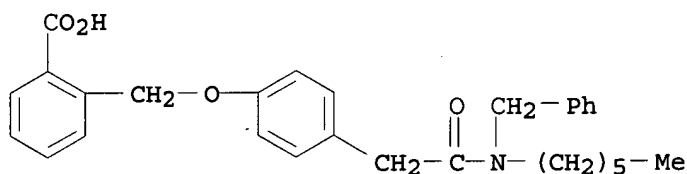
RN 637015-02-2 HCAPLUS

CN Benzoic acid, 2-[[[4-[2-[[2-(2,4-difluorophenyl)methyl]heptylamino]-2-oxoethyl]phenyl]thio]methyl]- (9CI) (CA INDEX NAME)



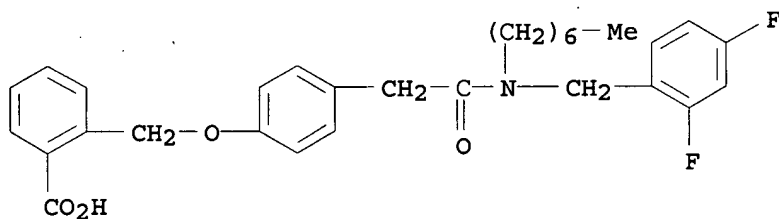
RN 637015-05-5 HCAPLUS

CN Benzoic acid, 2-[[[4-[2-[hexyl(phenylmethyl)amino]-2-oxoethyl]phenoxy]methyl]- (9CI) (CA INDEX NAME)



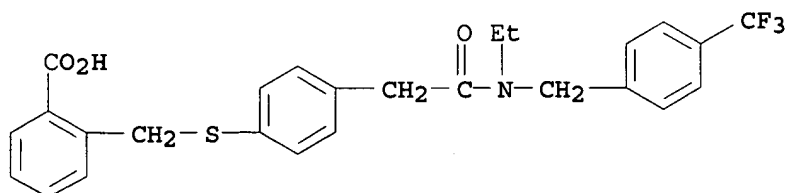
RN 637015-07-7 HCAPLUS

CN Benzoic acid, 2-[[[4-[2-[[2-(2,4-difluorophenyl)methyl]heptylamino]-2-oxoethyl]phenoxy]methyl]- (9CI) (CA INDEX NAME)



RN 637015-32-8 HCAPLUS

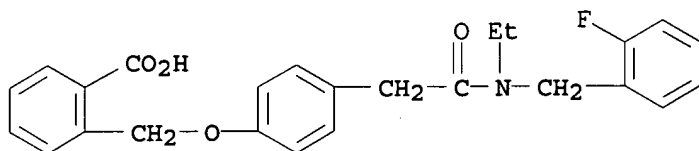
CN Benzoic acid, 2-[[[4-[2-[ethyl[[4-(trifluoromethyl)phenyl]methyl]amino]-2-oxoethyl]phenyl]thio]methyl]- (9CI) (CA INDEX NAME)



RN 637015-33-9 HCAPLUS

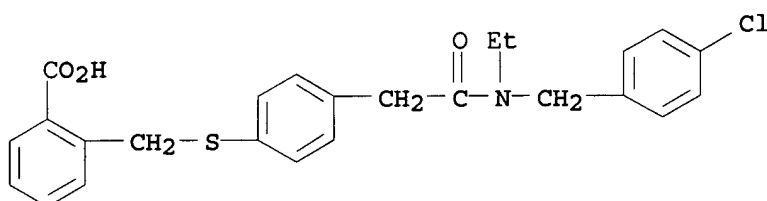
10/518,007

CN Benzoic acid, 2-[[4-[2-[ethyl[(2-fluorophenyl)methyl]amino]-2-oxoethyl]phenoxy]methyl]- (9CI) (CA INDEX NAME)



RN 637015-40-8 HCAPLUS

CN Benzoic acid, 2-[[[4-[2-[[[4-chlorophenyl)methyl]ethylamino]-2-oxoethyl]phenyl]thio]methyl]- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 2 OF 2 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2004:2679 HCAPLUS

DOCUMENT NUMBER: 140:76898

TITLE: Preparation of benzoic acid derivatives as modulators of PPAR- α and PPAR- γ

INVENTOR(S): Li, Lanna

PATENT ASSIGNEE(S): Astrazeneca AB, Swed.; Astrazeneca UK Limited

SOURCE: PCT Int. Appl., 101 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

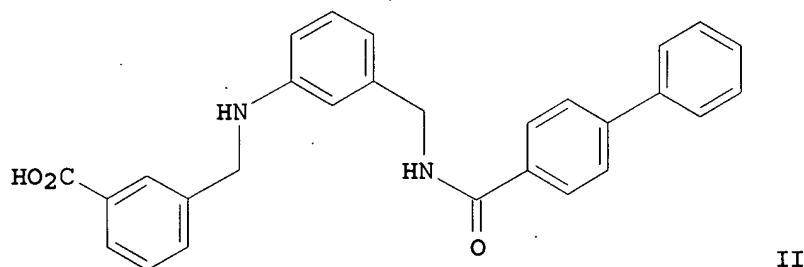
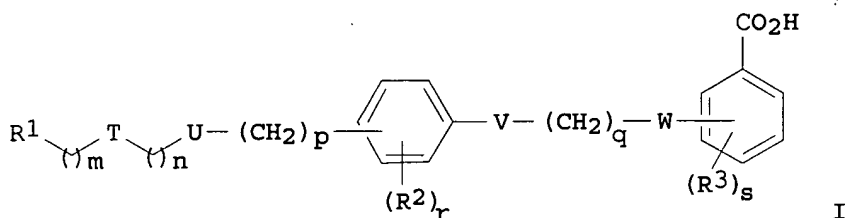
FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

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RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
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AU 2003240101	A1	20040106	AU 2003-240101	20030617
BR 2003011840	A	20050315	BR 2003-11840	20030617
EP 1517680	A1	20050330	EP 2003-732715	20030617
R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK			
CN 1662230	A	20050831	CN 2003-814319	20030617

10/518,007

JP 2006502105	T2	20060119	JP 2004-515010	20030617
NZ 536972	A	20060630	NZ 2003-536972	20030617
NO 2004005222	A	20050119	NO 2004-5222	20041129
ZA 2004009690	A	20051011	ZA 2004-9690	20041130
US 2005267149	A1	20051201	US 2004-518819	20041220
PRIORITY APPLN. INFO.:			SE 2002-1937	A 20020620
			WO 2003-GB2598	W 20030617
OTHER SOURCE(S):	MARPAT 140:76898			
GI				



AB Title compds. I [R1 = (un)substituted aryl, alkyl, acyl, etc.; (CH2)m-T-(CH2)n-U-(CH2)p = attached at either the meta or para position (to V) and is O(CH2)2, O(CH2)3, etc.; V = O, S, amino, single bond; q = 1-3; W = O, S, amido, amino, single bond; R2 = halo, alkyl, alkoxy, etc.; r = 0-3; R3 = halo, alkyl, alkoxy, etc.; s = 0-3; with some provisions] are prepared For instance, tert-Bu [3-[[[(1,1'-biphenyl-4-yl)carbonyl]amino]methyl]phenyl]carbamate (preparation given) is deprotected (CH2Cl2, TFA) and alkylated with 3-carboxybenzaldehyde (HOAc, NaBH4) to give II. Compds. of the invention have an EC50 < 50μmol/L for PPAR-α and PPAR-γ. I are useful in treating clin. conditions associated with insistance.

IT 637358-56-6P, 2-[[[4-[2-[Benzyl(hexyl)amino]-2-oxoethyl]-2-fluorophenoxy]methyl]benzoic acid 637358-59-9P, 2-[[[4-[2-[Benzyl(hexyl)amino]-2-oxoethyl]-2-methoxyphenoxy]methyl]benzoic acid

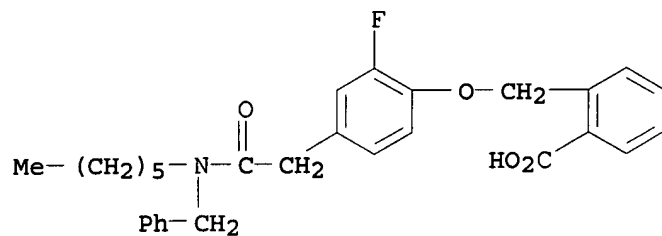
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of benzoic acid derivs. as modulators of PPAR-α and PPAR-γ)

RN 637358-56-6 HCAPLUS

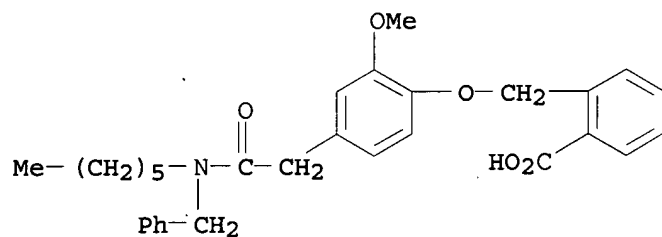
10/518,007

CN Benzoic acid, 2-[[2-fluoro-4-[2-[hexyl(phenylmethyl)amino]-2-oxoethyl]phenoxy]methyl]- (9CI) (CA INDEX NAME)



RN 637358-59-9 HCAPLUS

CN Benzoic acid, 2-[[4-[2-[hexyl(phenylmethyl)amino]-2-oxoethyl]-2-methoxyphenoxy]methyl]- (9CI) (CA INDEX NAME)



REFERENCE COUNT:

6

THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT